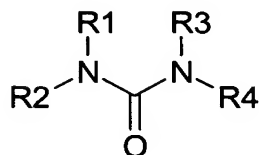


## Amendments to the Claims

Claim 1 (currently amended): A compound ~~which is represented by~~ comprising formula (I) ~~below~~



(I)

wherein

R<sub>1</sub> is CH<sub>3</sub> or CH<sub>2</sub>CH<sub>3</sub>;

R<sub>2</sub> is a *para* and/or *meta* substituted phenyl group;

R<sub>3</sub> is H, CH<sub>3</sub> or CH<sub>2</sub>CH<sub>3</sub>; and

R<sub>4</sub> is a linear or cyclic aliphatic group, ~~which is optionally substituted,~~

or, wherein

R<sub>1</sub> and R<sub>2</sub> are as stated above while R<sub>3</sub> and R<sub>4</sub> are parts of a 4- to 6-membered cyclic entity, ~~which is optionally substituted,~~

and which compound has affinity for human IgG of κ-type.

Claim 2 (currently amended): ~~A compound according to~~ The compound of claim 1, which is an affinity ligand with affinity for the constant region of a Fab fragment of human IgG of κ-type.

Claim 3 (currently amended): ~~A compound according to~~ The compound of claim 1 ~~or~~ 2, wherein R<sub>1</sub> is CH<sub>3</sub>.

Claim 4 (currently amended): ~~A~~ The compound of claim 1 according to any one of the preceding claims, wherein R<sub>2</sub> ~~comprises~~ is a substituted phenyl group ~~and the having~~ substituents ~~are selected from the group that consists~~ consisting of F, Cl, Br, I and O.

Claim 5 (currently amended): ~~A compound according to any one of the preceding claims~~, The compound of claim 1, wherein the phenyl group of R<sub>2</sub> is substituted in the *para* position with a group ~~defined as~~ -O-R<sub>5</sub>, wherein R<sub>5</sub> is either CH<sub>3</sub> or CH<sub>2</sub>CH<sub>3</sub>.

Claim 6 (currently amended): ~~A compound according to any one of the preceding claims~~, The compound of claim 4, wherein the phenyl group of R<sub>2</sub> is substituted with Cl or F in the *meta* position.

Claim 7 (currently amended): ~~A compound according to any one of claims 1-4~~, The compound of claim 4, wherein the phenyl group of R<sub>2</sub> is substituted with Cl in *meta* and *para* position.

Claim 8 (currently amended): ~~A compound according to any one of the preceding claims~~, The compound of claim 1, wherein R<sub>4</sub> is an aliphatic group, which includes oxygen atoms in one or more positions ~~is interrupted in one or more positions by oxygen atoms~~.

Claim 9 (currently amended): ~~A compound according to any one of the preceding claims~~, The compound of claim 1, wherein R<sub>4</sub> is an aliphatic group, which ~~comprises~~ contains one or more carbonyl ~~group groups~~.

Claim 10 (currently amended): ~~A compound according to any one of the preceding claims,~~ The compound of claim 1, wherein R<sub>4</sub> is an aliphatic group ~~that comprises~~ which includes a terminating functionality selected from the group ~~that consists~~ consisting of a carboxylic acid, nitrogen, oxygen, sulphur or any derivative thereof.

Claim 11 (currently amended): ~~A compound according to any one of the preceding claims,~~ The compound of claim 1, wherein R<sub>1</sub> is CH<sub>3</sub>; R<sub>2</sub> is a phenyl group that has been substituted with Cl in *meta* and *para* position; and R<sub>3</sub> and R<sub>4</sub> are parts of a cyclic 5-membered group, ~~which is optionally substituted.~~

Claim 12 (currently amended): ~~A compound according to~~ The compound of claim 11, wherein the cyclic 5-membered entity is substituted in a position directly adjacent to N with a C(O)-O-CH<sub>3</sub> group.

Claim 13 (currently amended): ~~A compound according to any one of the preceding claims,~~ The compound of claim 1, which is capable of binding ~~human~~ to the constant region of a human IgG of  $\kappa$ -type, or a functional derivative thereof, with a binding constant of at least 10<sup>-3</sup> M.

Claim 14 (currently amended): ~~A compound according to any one of the preceding claims,~~ The compound of claim 1, which is capable of binding to the constant region of a human IgG of  $\kappa$ -type, or a functional derivative thereof, via a binding pocket defined by the structure coordinates of the amino acids as shown in Fig 6.

Claim 15 (cancelled)

Claim 16 (currently amended): A sorption complex ~~comprised of a compound according to any one of claims 1-14~~ comprising the compound of claim 1 directly linked to the constant region of a Fab fragment of a human IgG of  $\kappa$ -type, or a functional derivative thereof.

Claim 17 (currently amended): A separation matrix for affinity chromatography, ~~which matrix comprises~~ comprising ligands coupled to a support, wherein the majority of the ligands are the compounds of claim 1 ~~as defined in any one of claims 1-14~~.

Claim 18 (currently amended): ~~A~~ The separation matrix ~~according to~~ of claim 17, wherein the ligands have been coupled to the support via linkers.

Claim 19 (currently amended): ~~A~~ The separation matrix ~~according to~~ of claim 17 ~~or 18~~, wherein the support is a porous polymeric particle.

Claim 20 (cancelled)

Claim 21 (currently amended): A system suitable for affinity chromatography, ~~which is comprised of a separation matrix as defined in any one of claims 17-19~~ comprising the separation matrix of claim 17 packed in a column.